#### § 1065.267

### § 1065.267 Gas chromatograph with a flame ionization detector.

- (a) Application. You may use a gas chromatograph with a flame ionization detector (GC-FID) to measure CH<sub>4</sub> concentrations of diluted exhaust for batch sampling. While you may also use a nonmethane cutter to measure CH<sub>4</sub>, as described in §1065.265, use a reference procedure based on a gas chromatograph for comparison with any proposed alternate measurement procedure under §1065.10.
- (b) Component requirements. We recommend that you use a GC-FID that meets the specifications in Table 1 of §1065.205 and that the measurement be done according to SAE J1151 (incorporated by reference in §1065.1010). The GC-FID must meet the linearity verification in §1065.307.

[76 FR 57442, Sept. 15, 2011, as amended at 79 FR 23761, Apr. 28, 2014]

## § 1065.269 Photoacoustic analyzer for ethanol and methanol.

- (a) Application. You may use a photoacoustic analyzer to measure ethanol and/or methanol concentrations in diluted exhaust for batch sampling.
- (b) Component requirements. We recommend that you use a photoacoustic analyzer that meets the specifications in Table 1 of §1065.205. Note that your photoacoustic system must meet the verification in §1065.369 and it must also meet the linearity verification in §1065.307. Use an optical wheel configuration that gives analytical priority to measurement of the least stable components in the sample. Select a sample integration time of at least 5 seconds. Take into account sample chamber and sample line volumes when determining flush times for your instrument.

[79 FR 23761, Apr. 28, 2014]

NO<sub>X</sub> AND N<sub>2</sub>O MEASUREMENTS

#### § 1065.270 Chemiluminescent detector.

(a) Application. You may use a chemiluminescent detector (CLD) to measure  $NO_X$  concentration in raw or diluted exhaust for batch or continuous sampling. We generally accept a CLD for  $NO_X$  measurement, even though it measures only NO and  $NO_2$ , when coupled with an  $NO_2$ -to-NO converter,

- since conventional engines aftertreatment systems do not emit significant amounts of NO<sub>X</sub> species other than NO and NO2. Measure other NO<sub>x</sub> species if required by the standard-setting part. While you may also use other instruments to measure NO<sub>X</sub>, as described in §1065.272, use a refprocedure based on erence chemiluminescent detector for comparison with any proposed alternate measurement procedure under §1065.10.
- (b) Component requirements. We recommend that you use a CLD that meets the specifications in Table 1 of §1065.205. Note that your CLD-based system must meet the quench verification in §1065.370 and it must also meet the linearity verification in §1065.307. You may use a heated or unheated CLD, and you may use a CLD that operates at atmospheric pressure or under a vacuum.
- (c)  $NO_2$ -to-NO converter. Place upstream of the CLD an internal or external  $NO_2$ -to-NO converter that meets the verification in §1065.378. Configure the converter with a bypass line if it is needed to facilitate this verification.
- (d) *Humidity effects*. You must maintain all CLD temperatures to prevent aqueous condensation. If you remove humidity from a sample upstream of a CLD, use one of the following configurations:
- (1) Connect a CLD downstream of any dryer or chiller that is downstream of an  $NO_2$ -to-NO converter that meets the verification in §1065.378.
- (2) Connect a CLD downstream of any dryer or thermal chiller that meets the verification in §1065.376.
- (e) Response time. You may use a heated CLD to improve CLD response time.

[70 FR 40516, July 13, 2005, as amended at 73 FR 37300, June 30, 2008; 76 FR 57442, Sept. 15, 2011; 79 FR 23761, Apr. 28, 2014]

# § 1065.272 Nondispersive ultraviolet analyzer.

(a) Application. You may use a non-dispersive ultraviolet (NDUV) analyzer to measure  $NO_X$  concentration in raw or diluted exhaust for batch or continuous sampling. We generally accept an NDUV for  $NO_X$  measurement, even though it measures only NO and  $NO_2$ , since conventional engines and

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aftertreatment systems do not emit significant amounts of other  $NO_X$  species. Measure other  $NO_X$  species if required by the standard-setting part. Note that good engineering judgment may preclude you from using an NDUV analyzer if sampled exhaust from test engines contains oil (or other contaminants) in sufficiently high concentrations to interfere with proper operation.

- (b) Component requirements. We recommend that you use an NDUV analyzer that meets the specifications in Table 1 of §1065.205. Note that your NDUV-based system must meet the verifications in §1065.372 and it must also meet the linearity verification in §1065.307.
- (c) NO<sub>2</sub>-to-NO converter. If your NDUV analyzer measures only NO, place upstream of the NDUV analyzer an internal or external NO<sub>2</sub>-to-NO converter that meets the verification in §1065.378. Configure the converter with a bypass to facilitate this verification.
- (d) *Humidity effects*. You must maintain NDUV temperature to prevent aqueous condensation, unless you use one of the following configurations:
- (1) Connect an NDUV downstream of any dryer or chiller that is downstream of an  $NO_2$ -to-NO converter that meets the verification in §1065.378.
- (2) Connect an NDUV downstream of any dryer or thermal chiller that meets the verification in §1065.376.

[70 FR 40516, July 13, 2005, as amended at 73 FR 59323, Oct. 8, 2008; 76 FR 57442, Sept. 15, 2011; 79 FR 23761, Apr. 28, 2014]

#### § 1065.275 N<sub>2</sub>O measurement devices.

- (a) General component requirements. We recommend that you use an analyzer that meets the specifications in Table 1 of §1065.205. Note that your system must meet the linearity verification in §1065.307.
- (b) *Instrument types*. You may use any of the following analyzers to measure N<sub>2</sub>O:
- (1) Nondispersive infrared (NDIR) analyzer.
- (2) Fourier transform infrared (FTIR) analyzer. Use appropriate analytical procedures for interpretation of infrared spectra. For example, EPA Test Method 320 is considered a valid method for spectral interpretation (see

http://www.epa.gov/ttn/emc/methods/method320.html).

- (3) Laser infrared analyzer. Examples of laser infrared analyzers are pulsed-mode high-resolution narrow band midinfrared analyzers, and modulated continuous wave high-resolution narrow band mid-infrared analyzers.
- (4) Photoacoustic analyzer. Use an optical wheel configuration that gives analytical priority to measurement of the least stable components in the sample. Select a sample integration time of at least 5 seconds. Take into account sample chamber and sample line volumes when determining flush times for your instrument.
- (5) Gas chromatograph analyzer. You may use a gas chromatograph with an electron-capture detector (GC–ECD) to measure  $N_2O$  concentrations of diluted exhaust for batch sampling.
- (i)You may use a packed or porous layer open tubular (PLOT) column phase of suitable polarity and length to achieve adequate resolution of the  $N_2O$  peak for analysis. Examples of acceptable columns are a PLOT column consisting of bonded polystyrenedivinylbenzene or a Porapack Q packed column. Take the column temperature profile and carrier gas selection into consideration when setting up your method to achieve adequate  $N_2O$  peak resolution.
- (ii) Use good engineering judgment to zero your instrument and correct for drift. You do not need to follow the specific procedures in §\$1065.530 and 1065.550(b) that would otherwise apply. For example, you may perform a span gas measurement before and after sample analysis without zeroing and use the average area counts of the pre-span and post-span measurements to generate a response factor (area counts/span gas concentration), which you then multiply by the area counts from your sample to generate the sample concentration.
- (c) Interference verification. Perform interference verification for NDIR, FTIR, laser infrared analyzers, and photoacoustic analyzers using the procedures of \$1065.375. Interference verification is not required for GC-ECD. Certain interference gases can positively interfere with NDIR, FTIR, and photoacoustic analyzers by causing